INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference 12373580/JGC	FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416).					
Internat: aal Application No.	International Filing Dat (day/month/year)	te Priority Date (day/month/year)					
PCT/AU29#3/001606	2 December 2003	2 December 2002	_				
International Patent Classification (IPC) or r	International Patent Classification (IPC) or national classification and IPC						
Int. Cl. 7 C07K 7/06, 7/08: A61K 38/0	04; A61P 9/00, 13/00,	25/00					
Applicant XENOME LTD et al							
This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.							
2. This REPORT consists of a total of 3	_						
This report is also accompanied by ANNEXES, i.e., sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).							
These annexes consist of a total of	of 2 sheet(s).						
3. This report contains indications relating	g to the following items:						
I X Basis of the report							
II Priority		•	!				
III Non-establishment of op	inion with regard to nov	velty, inventive step and industrial applicability					
IV Lack of unity of inventio	on .	·					
V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement							
VI Certain documents cited	VI Certain documents cited						
VII Certain defects in the int	VII Certain defects in the international application						
VIII Certain observations on the international application							
Date of submission of the demand Date of completion of the report							
25 March 2004		11 June 2004					
Name and mailing address of the IPEA/AU		Authorized Officer					
AUSTRALIAN PATENT OFFICE PO BOX 200, WODEN ACT 2606, AUSTRALIA							
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I.		is of the report					
1.		regard to the elements of the international application:*					
	=	e international a	application as originally filed.				
	X the	e description,	pages 1, 3-30, 32-59, as originally filed,				
			pages , filed with the demand,				
			pages 2, 31, received on 1 June 2004 with the letter of 1 June 2004				
	X the	e claims,	pages 60-67, as originally filed,				
			pages, as amended (together with any statement) under Article 19,				
	•		pages , filed with the demand,				
	· ·		pages, received on with the letter of				
	X the	drawings,	pages 1/1, as originally filed,				
		·	pages, filed with the demand, pages, received on with the letter of				
	X the	e sequence listi	ng part of the description:				
	<u>V</u>	o sequence mon	pages 1-84, as originally filed				
			pages, filed with the demand				
	-		pages, received on with the letter of				
2.			uage, all the elements marked above were available or furnished to this Authority in the language in				
		hich the international application was filed, unless otherwise indicated under this item. These elements were available or furnished to this Authority in the following language which is:					
•	the	the language of a translation furnished for the purposes of international search (under Rule 23.1(b)).					
			ublication of the international application (under Rule 48.3(b)).				
		e language of the d/or 55.3).	ne translation furnished for the purposes of international preliminary examination (under Rules 55.2				
3.		regard to any nucleotide and/or amino acid sequence disclosed in the international application, the international eliminary examination was carried out on the basis of the sequence listing:					
			nternational application in written form.				
	Fil	Filed together with the international application in computer readable form.					
	fur	furnished subsequently to this Authority in written form.					
	fur	furnished subsequently to this Authority in computer readable form.					
The statement that the subsequently furnished written sequence listing does not go beyond the disclosure international application as filed has been furnished.							
		ne statement tha en furnished	at the information recorded in computer readable form is identical to the written sequence listing has				
4.	Tb	e amendments	have resulted in the cancellation of:				
		the descr	ription, pages				
		the claim	ns, Nos.				
•		the draw	ings, sheets/fig.				
5.			een established as if (some of) the amendments had not been made, since they have been considered to colosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).**				
*		Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17).					
**	Any rep	Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report					

V.	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations
	and explanations supporting such statement

1.	Statement				
	Novelty (N)	Claims 1-37	YES		
	•	Claims	МО		
	Inventive step (IS)	Claims 1-37	YES .		
		Claims	МО		
	Industrial applicability (IA)	Claims 1-37	YES		
		Claims	NO		

2. Citations and explanations (Rule 70.7)

The following documents were cited in the International Search Report:

D1: WO 2000/020444 D2: WO 2000/044769

Claims 1-37 are novel because neither D1 or D2 explicitly disclose χ -conotoxins with peptides with sequences corresponding to SEQ ID NO 5 or 6, and in the case of SEQ ID NO 3 and 4, peptides χ -MrIA, χ -MrIB, Mar2 have been excluded from the scope of the claims as a result of the proviso. Mar 1 of D2 does fall within the scope of SEQ ID NO 3 and 4, but is not disclosed as a χ -conotoxin peptide.

D1 discloses the χ -conotoxin peptides χ -MrIA and χ -MrIB which are of close sequence homology to SEQ ID NO 5 and 6 of the current invention, and furthermore contemplates derivatives, including addition or substitution of amino acids. However D1 has not disclosed the specific modifications as in the current claims, and given the applicant has found certain unexpected advantages of SEQ ID NO 5 over χ -MrIA (see page 7 lines 22-31 of the current application) and that the derivative according to SEQ ID NO 6 increases the binding affinity of the χ -peptides, it can be acknowledged that claims 1-37 are inventive when compared to D1.

D2 discloses peptides which appear to be α -conotoxins, in which case D2 teaches away from the χ -conotoxins of the current application. D2 discloses Mar1 and Mar2 which are also of close sequence homology to SEQ ID NO 5 and 6, but, again they are α -conotoxins, not χ -conotoxins. Even if they were χ -conotoxins, by using the same reasoning as for D1, given the unexpected advantages the applicant has discovered for these polypeptides claims 1-37 are inventive when compared to D2.

Claims 1-37 meet the requirement for industrial applicability.